CLAIMS

We claim:

1. A compound of a compound of formula I:

its enantiomeric, diastereomeric, or tautomeric isomer thereof, or a pharmaceutically acceptable salt thereof wherein,

G is phenyl substituted with from one (1) to five (5) R¹ substituents;

each R¹ is independently

- 15 (a) Cl,
 - (b) Br,
 - (c) F,
 - (d) CN,
 - (e) C_{1-7} alkyl, or
- 20 (f) NO₂;

R² is

25

30

- (a) H,
- (b) R^5 ,
- (c) NR^7R^8 ,
- (d) SO_2R^{10} , or
 - (e) OR^9 ;

A is C_{1-7} alkyl;

W is a five- (5) or six- (6) membered heterocyclic ring having one (1), two (2) or three (3) heteroatoms selected from the group consisting of O, $S(O)_k$, and N wherein W is optionally substituted with one or more OH, oxo (=O), or C_{1-7} alkyl;

B is

- (a) C₁₋₇alkyl optionally substituted by OH or NR⁷R⁸,
- (b) O, or

(c) NR^{11} ;

 R^3 is

5

10

15

- (a) phenyl, optionally fused to a benzene or pyridine ring, and optionally substituted by R¹², wherein optionally any two adjacent R¹² substituents taken together constitute a group of the formula -O(CH₂)O-,
 -(NH)C(=O)(CH₂)_iO-, or -(CH₂)_i-, or
- (b) a five- (5) or six- (6) membered heteroaryl bonded via a carbon atom having one (1), two (2), or three (3) heteroatoms selected from the group consisting of O, S, and N-Z, wherein R³ is optionally fused to a benzene or pyridine ring, and optionally substituted with one or more R¹², wherein Z is absence, H, or C₁₋₄alkyl;

R4 is

- (a) H,
- (b) halo, or
- (c) C₁₋₄alkyl optionally substituted by halo;

R⁵ is

- (a) $(CH_2)_mOCH_2CH_2OR^{11}$,
- (b) het, wherein said het is bound via a carbon atom,
- (c) aryl,
- 20 (d) C₁₋₇alkyl which may be partially unsaturated and is optionally substituted by one or more R⁶ substituents, or
 - (e) C_{3-8} cycloalkyl which may be partially unsaturated and optionally substituted by one or more R^6 or C_{1-7} alkyl optionally substituted by R^6 ;

R⁶ is

25 (a) OR^9 ,

- (b) SR^9 ,
- (c) NR^7R^8 ,
- (d) halo,
- (e) $CONR^7R^8$,
- 30 (f) CO_2R^9 ,
 - (g) het,
 - (h) phenyl, optionally substituted by R¹²,
 - (i) CN,

- (j) oxo,
- (k) $SO_2NR^9R^{11}$,
- (1) SO_mR^{10} , or
- (m) $P(=O)(OR^{11})(R^{11});$
- 5 R⁷ and R⁸ are independently
 - (a) H,
 - (b) aryl,
 - (c) C₁₋₇alkyl which may be partially unsaturated and is optionally substituted by one or more NR¹¹R¹¹, OR¹¹, SR¹¹, SO_mR¹⁰, CONR¹¹R¹¹, CO₂R¹¹, het, aryl, cyano, or halo,
 - (d) C₃₋₈cycloalkyl,
 - (e) $(C=O)R^{10}$, or
 - (f) R⁷ and R⁸ together with the nitrogen to which they are attached form a het;
- 15 R^9 is

10

- (a) H,
- (b) aryl,
- (c) het, wherein the het is bound through a carbon atom,
- (d) C₁₋₇alkyl which is optionally partially unsaturated and is optionally substituted by one or more aryl, het, OR¹¹, SR¹¹, NR¹¹R¹¹, halo, or C₃₋₈cycloalkyl substituents and which C₃₋₈cycloalkyl is optionally substituted by OR¹¹, or
 - (e) C₃₋₈cycloalkyl which is optionally partially unsaturated and is optionally substituted by one or more halo, OR¹¹, SR¹¹, or NR¹¹R¹¹ substituents;
- 25 R¹⁰ is

- (a) aryl,
- (b) het,
- (c) C₁₋₇alkyl which is optionally partially unsaturated and is optionally substituted by one or more aryl, het, OR¹¹, SR¹¹, NR¹¹R¹¹, halo, or C₃₋₈cycloalkyl substituents and which C₃₋₈cycloalkyl is optionally substituted by OR¹¹, or
 - (d) C₃₋₈cycloalkyl which is optionally partially unsaturated and is optionally substituted by one or more halo, OR¹¹, SR¹¹, or NR¹¹R¹¹ substituents;

R¹¹ is (a) H, or (b) C₁₋₇alkyl; R¹² is halo, 5 (a) OR14, (b) SR¹¹, (c) NR^7R^8 , (d) (e) phenyl, optionally substituted by halo, C₁₋₇alkyl, or C₁₋₇alkoxy, C₁₋₇alkyl which is optionally partially unsaturated and optionally **(f)** 10 substituted by R¹³, cyano, (g) (h) nitro, CONR⁷R⁸, (i) SO₂NR⁷R⁸, (j) 15 CO₂R¹¹, or (k) · $NHC(=O)R^{11}$; **(1)** R¹³ is (a) phenyl, optionally substituted by halo, C₁₋₇alkyl, or C₁₋₇alkoxy, OR11, 20 (b) $O(CH_2CH_2O)_nR^{11}$, (c) NR⁷R⁸, or (d) (e) halo; R¹⁴ is (a) Η 25 alkyl, optionally substituted by halo, (b) phenyl, optionally substituted by halo, C₁₋₇alkyl, or C₁₋₇alkoxy, or (c)

wherein any aryl is optionally substituted with one or more substituents selected from the group consisting of halo, OR¹¹, NR¹¹R¹¹, cyano, CO₂R¹¹, or C₁₋₇alkyl in which said C₁₋₇alkyl is optionally substituted by one to three halo, OR¹¹, or NR¹¹R¹¹;

 $-(CH_2CH_2O)_nOR^{11};$

(d)

wherein any het is optionally substituted with one or more substituents selected from the group consisting of halo, OR^{11} , $NR^{11}R^{11}$, cyano, CO_2R^{11} , oxo (=0), or $C_{1.7}$ alkyl in which said $C_{1.7}$ alkyl is optionally substituted by one to three halo, OR^{11} , or $NR^{11}R^{11}$;

```
i is 3 or 4;
j is 0 or 1;
k is 0, 1, or 2;
each n is independently 1, 2, 3, 4 or 5; and
each m is independently 1 or 2;
```

10

- 2. A compound of claim 1 wherein R¹ is F, Cl, or cyano.
- 3. A compound of claim 2 wherein R^1 is Cl.

15

- 4. A compound of claim 2 wherein R^1 is F.
- 5. A compound of claim 1 wherein G is 4-chlorophenyl.
- 20 6. A compound of claim 1 wherein G is 4-fluorophenyl.
 - 7. A compound of claim 1 wherein R² is H.
 - 8. A compound of claim 1 wherein R^2 is R^5 .

- 9. A compound of claim 1 wherein R² is NR⁷R⁸.
- 10. A compound of claim 1 wherein R² is SO₂R¹⁰.
- 30 11. A compound of claim 1 wherein R² is OR⁹.
 - 12. A compound of claim 8 wherein R^2 is C_{1-7} alkyl which may be partially unsaturated and is optionally substituted with one or more R^6 substituents.

13.	Α	compound	of	claim	12	wherein	R^2	is	methy	vl.
x		O LIAP O GALGO	\sim $_{\rm L}$	ATCOURTE		******		-		,

- 14. A compound of claim 12 wherein R² is ethyl.
- 15. A compound of claim 1 wherein A is C₁₋₄alkyl.
- 16. A compound of claim 1 wherein A is methyl.
- 17. A compound of claim 1 wherein W is a six- (6) membered heterocyclic ring having one (1), two (2), or three (3) heteroatoms selected from the group consisting of O, S(O)_k, or N, wherein het is optionally substituted with C₁₋₄ alkyl.
- 18. A compound of claim 1 wherein W is a five- (5) membered heterocyclic ring having one (1), two (2), or three (3) heteroatoms selected from the group consisting of O, S(O)_k, or N, wherein het is optionally substituted with C₁₋₄ alkyl.
- 20 19. A compound of claim 17 wherein W is morpholine.
 - 20. A compound of claim 18 wherein W is pyrrolidine.
 - 21. A compound of claim 1 wherein B is C₁₋₄alkyl.
 - 22. A compound of claim 1 wherein B is methyl.
 - 23. A compound of claim 1 wherein B is methyl substituted with a hydroxy.
- 30 24. A compound of claim 1 wherein R³ is phenyl.
 - 25. A compound of claim 1 wherein R³ is naphthyl

00680.US1

5

10

- 26. A compound of claim 1 wherein R³ is phenyl, fused to a pyridine ring.
- 27. A compound of claim 1 wherein R³ is a five- (5) membered heteroaryl bonded via a carbon atom having one (1) or two (2) heteroatoms selected from the group consisting of O, S, and N-Z.
- 28. A compound of claim 1 wherein R³ is a five- (5) membered heteroaryl bonded via a carbon atom having one (1) or two (2) heteroatoms selected from the group consisting of O, S, and N-Z, wherein R³ is fused to a benzene or pyridine ring.
- 29. A compound of claim 1 wherein R³ is a six- (6) membered heteroaryl bonded via a carbon atom having one (1) or two (2) nitrogen atoms.
- 15 30. A compound of claim 1 wherein R³ is a six- (6) membered heteroaryl bonded via a carbon atom having one (1) nitrogen atom.
- 31. A compound of claim 1 wherein R³ is a six- (6) membered heteroaryl bonded via a carbon atom having one (1) or two (2) nitrogen atoms and is fused to a benzene ring.
 - 32. A compound as in any of claims 24 31 wherein R^3 is substituted by R^{12} .
- 33. A compound of claim 27 wherein R³ is 2-furyl, thien-2-yl, 1,3-thiazol-2-yl, 1,3-thiazol-5-yl, or 1H-imidazol-2-yl.
 - 34. A compound of claim 29 wherein R³ is pyrimidin-2-yl, or pyrimidin-5-yl.
 - 35. A compound of claim 29 wherein R³ is pyrazin-2-yl.
 - 36. A compound of claim 30 wherein R³ is pyridin-2-yl, or pyridin-3-yl.

- 37. A compound of claim 1 wherein R³ is 1,3-benzoxazol-2-yl, or 1,3-benzothiazol-2-yl.
- 38. A pharmaceutical composition comprising a compound of claim 1 and a pharmaceutically acceptable carrier.
 - 39. A method of treating infections by herpesviruses which comprises administering to a mammal in need thereof a compound of claim 1.
- 10 40. The method of claim 39 wherein said herpesviruses is herpes simplex virus types 1, herpes simplex virus types 2, varicella zoster virus, human cytomegalovirus, Epstein-Barr virus, human herpes virus 6, human herpes virus 7 or human herpes virus 8.
- 15 41. The method of claim 40 wherein said herpesviruses is human cytomegalovirus.
 - 42. The method of claim 40 wherein said herpesviruses is varicella zoster virus or Epstein-Barr virus.
- 20 43. The method of claim 40 wherein said herpesviruses is herpes simplex virus types 1 or herpes simplex virus types 2.
 - 44. The method of claim 39 wherein the compound of claim 1 is administered orally, parenterally or topically.
 - 45. The method of claim 39 wherein the compound of claim 1 is in an amount of from about 0.1 to about 300 mg/kg of body weight.
- 46. The method of claim 39 wherein the compound of claim 1 is in an amount of from about 1 to about 30 mg/kg of body weight.
 - 47. The method of claim 39 wherein said mammal is a human.

- 48. The method of claim 39 wherein said mammal is an animal.
- 49. A method of treating atherosclerosis and restenosis comprising administering to a mammal in need thereof a compound of claim 1.

15

20

- 45. A method for inhibiting a herpesviral DNA polymerase, comprising contacting the polymerase with an effective inhibitory amount of a compound of claim 1.
- 51. A compound of claim 1, or a pharmaceutically acceptable salt thereof, for use in the manufacture of medicines for the treatment or prevention of a herpesviral infection in a mammal.
 - 52. A compound of claim 1 which is
 - (1) 2-(((3S)-3-benzylmorpholin-4-yl)methyl)-N-(4-chlorobenzyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (2) N-(4-chlorobenzyl)-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide
 - (3) N-(4-Chlorobenzyl)-2-(((2R*)-2-((R*)-hydroxy(pyridin-2-yl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (4) N-(4-Chlorobenzyl)-2-(((2R*)-2-((R*)-2-furyl(hydroxy)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
- 25 (5) N-(4-chlorobenzyl)-2-(((2R)-2-((R)-hydroxy(1,3-thiazol-2-yl)methyl)-pyrrolidin-1-yl}methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (6) N-(4-chlorobenzyl)-2-(((2R)-2-((R)-hydroxy(thien-2-yl)methyl)-pyrrolidin-1-yl}methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,
 - (7) 2-(((2R)-2-((R)-1,3-benzothiazol-2-yl(hydroxy)methyl)pyrrolidin-1-yl)-methyl)-N-(4-chlorobenzyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,

15

20

(8)	N-(4-chlorobenzyl)-2-(((2 R)-2-((R)-hydroxy(1,3-thiazol-5-yl)methyl)-
pyrroli	idin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-
carbox	zamide.

- (9) N-(4-chlorobenzyl)-2-(((2R)-2-((R)-hydroxy(pyridin-2-yl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
- (10) N-(4-chlorobenzyl)-2-(((2R)-2-((S)-hydroxy(pyridin-3-yl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,
- 10 (11) N-(4-chlorobenzyl)-2-(((2R)-2-((S)-hydroxy(pyrimidin-5-yl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (12) N-(4-chlorobenzyl)-2-(((2R)-2-((R)-hydroxy(1H-imidazol-2-yl)-methyl)pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (13) 2-(((2R)-2-((R)-1,3-benzoxazol-2-yl(hydroxy)methyl)pyrrolidin-1-yl)-methyl)-N-(4-chlorobenzyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (14) N-(4-chlorobenzyl)-2-(((3R)-3-((R)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,
 - (15) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,
 - (16) N-(4-chlorobenzyl)-7-ethyl-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (17) N-(4-chlorobenzyl)-7-ethyl-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)-morpholin-4-yl)methyl)-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
- (18) N-(4-chlorobenzyl)-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)pyrrolidin-1-yl)methyl)-4-oxo-7-propyl-4,7-dihydrothieno[2,3-b]pyridine-5carboxamide,
 - (19) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-4-oxo-7-propyl-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,

15

25

- (20) N-(4-chlorobenzyl)-2-((($2R^*$)-2-((S^*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-7-(2-methoxyethyl)-4-oxo-4,7-dihydrothieno[2,3-b]-pyridine-5-carboxamide,
- (21) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-7-(2-methoxyethyl)-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (22) N-(4-chlorobenzyl)-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-7-(2-morpholin-4-ylethyl)-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
- 10 (23) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-7-(2-morpholin-4-ylethyl)-4-oxo-4,7-dihydrothieno[2,3-b]-pyridine-5-carboxamide,
 - (24) N-(4-chlorobenzyl)-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-3,7-dimethyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (25) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-3,7-dimethyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide, or a pharmaceutically acceptable salt thereof.
- 20 53. A compound of claim 1 which is
 - (1) 2-(((3S)-3-benzylmorpholin-4-yl)methyl)-N-(4-chlorobenzyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (2) N-(4-chlorobenzyl)-2-(((2R*)-2-((S*)-hydroxy(phenyl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide
 - (3) N-(4-Chlorobenzyl)-2-(((2R*)-2-((R*)-hydroxy(pyridin-2-yl)methyl)-pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide,
 - (4) N-(4-chlorobenzyl)-2-(((3R)-3-((R)-hydroxy(phenyl)methyl)morpholin-4-yl) methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide,
 - (5) N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy(phenyl)methyl)morpholin-4-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-B]pyridine-5-carboxamide, or a pharmaceutically acceptable salt thereof.

- 54. A compound of claim 1 which is *N*-(4-chlorobenzyl)-2-(((2*R**)-2-((*S**)-hydroxy(phenyl)methyl)pyrrolidin-1-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-*b*]pyridine-5-carboxamide, or a pharmaceutically acceptable salt thereof.
- 55. A compound of claim 1 which is N-(4-chlorobenzyl)-2-(((3R)-3-((S)-hydroxy-(phenyl)methyl)morpholin-4-yl)methyl)-7-methyl-4-oxo-4,7-dihydrothieno[2,3-b]pyridine-5-carboxamide, or a pharmaceutically acceptable salt thereof.